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10588818 - GAU: 1614

U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

IDS FORM INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Application Number	10/588,818
				Filed	August 9, 2006
				First Named Inventor	Takahide NISHI
				Group Art Unit	1614
				Examiner Name	Alicia R. HUGHES Nelson Blakely III
Sheet	1	of	15	Attorney Docket Number	06439/HG

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*English-language abstract enclosed; also JP 11-80026 is a related family member of USP 6,004,565; USP 6,667,025 and US 2004/0092603

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U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

IDS FORM INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Application Number	10/588,818
				Filed	August 9, 2006
				First Named Inventor	Takahide NISHI
				Group Art Unit	1614
				Examiner Name	Alicia R. HUGHES Nelson Blakely III
Sheet	11	of	15	Attorney Docket Number	06439/HG

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		<p>CATIVIELA et al., "Stereoselective synthesis of quaternary α-amino acids, Part 1: Acyclic compounds", <u>Tetrahedron: Asymmetry</u>, <u>9</u>, pp. 3517-3599 (1998).</p> <p>GANDER-COQUOZ et al., "Synthesis of Enantiomerically Pure, α-Alkylated Lysine, Ornithine, and Tryptophan Derivatives", <u>Helvetica Chimica Acta</u>, <u>71</u>, pp. 224-236, (1988).</p> <p>SANO et al., "Lewis Acid- and Cationic Lithium-Mediated Diastereoselective Aldol-Type Reaction Based on a Double Chiral Recognition Manner for the Asymmetric Synthesis of α-Substituted Serines", <u>Tetrahedron Letters</u>, <u>36</u>, No. 23, pp. 4101-4104 (1995)</p> <p>NAGAO et al., "Efficient Preparation of New Chiral Synthons Useful for (+)-Carbacyclin Synthesis by Utilizing Enzymatic Hydrolysis", <u>Chemistry Letters</u>, pp. 239-242 (1989).</p> <p>TAMAI et al., "Enzymatic Hydrolyses of the δ-Symmetric Dicarboxylic Diesters Bearing a Sulfinyl Group as the Prochiral Center", <u>Chemistry Letters</u>, pp. 2381-2384 (1994).</p> <p>CASARRUBIO et al., "On the Syntheses of Thiophene Analogs of Practolol and 'Reversed' Practolol", <u>J. Heterocyclic Chem.</u>, <u>20</u>, 1557-1560 (1983).</p> <p>CHARETTE et al., "Syntheses of α, α-Disubstituted-α-Amino Acids by Double Nucleophilic Addition to Cyanohydrins", <u>Tetrahedron Letters</u>, <u>39</u>, 5147-5150 (1998).</p>			
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		<p>HIROSE et al., "2-Aminoalcohol Immunosuppressants: Structure-Activity Relationships," <u>Bioorganic & Medicinal Chemistry Letters</u>, Vol. 6, No. 22, pp. 2647-2650, 1996.</p> <p>KLEY et al., "Synthesis and PLA₂-Inhibitory Properties of 2(R)-Acetamido-Alkylphosphomethanols with a Variable Aggregate Anchor," <u>Bioorganic & Medicinal Chemistry Letters</u>, 9 (1999) 261-264.</p> <p>MAIER et al., "Organic Phosphorus Compounds 93. Preparation, Properties and Herbicidal Activity of 2-Substituted 5-Phenoxy- and 5-Pyridyloxy-Phenylaminoalkyl-Phosphonic- and -Phosphinic Acid - as well as - Phosphine Oxide Derivatives," <u>Phosphorus, Sulfur and Silicon</u>, 56 (1-4): 5-15 (1991).</p> <p>U.S. Application No. 11/922,429 deposited December 18, 2007, Confirmation No. 6507.</p>			
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